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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/936,957	01/09/2002	Peter John Mcikle	016994-01401US	2903
7590	12/14/2004		EXAMINER	
Jackson Walker LLP 2435 N. Central Expressway suite 600 Richardson, TX 75080			LAM, ANN Y	
			ART UNIT	PAPER NUMBER
			1641	

DATE MAILED: 12/14/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary	Application No.	Applicant(s)	
	09/936,957	MEIKLE ET AL.	
	Examiner	Art Unit	
	Ann Y. Lam	1641	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --
Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) Responsive to communication(s) filed on 20 September 2004.
 2a) This action is **FINAL**. 2b) This action is non-final.
 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) Claim(s) 1-20 and 36 is/are pending in the application.
 4a) Of the above claim(s) 21-35,37 and 38 is/are withdrawn from consideration.
 5) Claim(s) _____ is/are allowed.
 6) Claim(s) 1-20 and 36 is/are rejected.
 7) Claim(s) _____ is/are objected to.
 8) Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) The specification is objected to by the Examiner.
 10) The drawing(s) filed on _____ is/are: a) accepted or b) objected to by the Examiner.
 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
 a) All b) Some * c) None of:
 1. Certified copies of the priority documents have been received.
 2. Certified copies of the priority documents have been received in Application No. _____.
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)
Paper No(s)/Mail Date _____ | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152) |
| | 6) <input type="checkbox"/> Other: _____ |

DETAILED ACTION

Claim Rejections - 35 USC § 112

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claim 18 is rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement. The claim(s) contains subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention. Claim 18, line 3, recites the limitation "a combination thereof". There is not support in the original specification for a detection of a combination of lysosomal storage disorder. (The specification discloses that an elevated level of saposin correlates with the presence of several lysosomal storage disorders (page 6, lines 8-9). However, this disclosure does not describe more than one lysosomal storage disorder being diagnosed in a single assay.)

Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 1-20 and 36 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 1 is vague because it is not clear as to what level of saposin indicates the presence of the lysosomal storage disorder. The claim is also not clear as to what levels of saposin are involved in the monitoring of the lysosomal storage disorder. The claim is also not clear as to what level of saposin and/or which saposin is correlated to each of the lysosomal storage disorders. Page 4 of the specification indicates there are 30 lysosomal disorders and page 5 of the specification indicates there are 4 saposins. The detection of which saposin and at which level is indicative of the presence of which of the 30 possible lysosomal disorders? Applicant has not indicated which specific lysosomal disorder, nor a specific level of saposin or a specific level of increase or specific level of decrease level of saposin indicates the presence of the specific disease. In line 6, the recitation of "similar or different" is vague and indefinite as to what level of saposin is determined.

Claim 1 is further vague, specifically parts (i) and (ii). If the first level is similar to the baseline level of the control population of patients unaffected by lysosomal storage disorder, how can the first level be an indicator of the presence or extent of the lysosomal storage disorder?

Claim 7 is also indefinite since it is not clear as to what the measured level that is greater than the 95% level in the control population indicates. For example, does it indicate the presence of the disorder?

Claims 11 and 12 are vague because it is not clear as to what level of saposin, or a specific level of increase or decrease of saposin, indicates progression of the disorder. The claim is also not clear as to which of the 30 possible lysosomal disorders is being referred to relative to the 4 possible saposins that are being measured.

Claim 13 is vague because the recitation of "the second saposin" lacks antecedent support. The claim is further vague because it does not further limit claim 1. Claim 1 already recites the saposins in claim 13.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

1. Claims 1-4, 7, 8, 13-15, 17 and 18 rejected under 35 U.S.C. 103(a) as being unpatentable over O'Brien et al. (1991) Saposin proteins: structure, function, and role in human lysosomal disorders, THE FASEB JOURNAL, vol. 5(3), 301-8, in view of Sano et al., "Sphingolipid hydrolase activator proteins and their precursors", Biochemical and biophysical research communications, 165 (3), pp. 1191-7, 1989.

O'Brien teaches the invention substantially as claimed. More specifically, as to claim 1, O'Brien discloses the method of monitoring a lysosomal storage disorder in a

patient (page 306, right column, lines 21-22), comprising: measuring the level of at least one saposin in a tissue sample of the patient (page 306, right column, lines 17-19), wherein the level is an indicator of presence or extent of the disorder in the patient (page 306, right column, second full paragraph.)

As to claim 4, the measured level exceeds a mean level in a control population of individuals not having a lysosomal storage disorder, to indicate presence of the disorder in a patient (page 306, right column, lines 41-42.)

As to claim 7, the measured level is greater than the 95% level in the control population (page 306, right column, lines 41-42.)

As to claim 8, the patient is not known to have a lysosomal storage disorder before the measuring step (page 306, second full paragraph.)

As to claims 13 and 14, the saposin is selected from the group consisting of saposin A, B, C, D, and prosaposin (for example saposin A, page 306, right column, line 41.)

As to claim 15, the measuring step comprises detecting binding between a saposin polypeptide and an antibody (page 306, left column, lines 10-11.)

As to claim 17, the antibody is immobilized to a solid phase (page 306, right column, , line 18.)

As to claim 18, the lysosomal storage disorder is Niemann-Pick disease (page 306, right column, line 61.)

O'Brien teaches the detection of saposin, and its deficiency or accumulation, in specific tissue and cell samples as an indication of lysosomal storage disorder.

However, O'Brien does not specifically teach detection of saposin in whole blood or plasma samples (claims 1-3.)

Sano teaches that saposin is found in human blood and plasma (see abstract.) It would have been obvious to one of ordinary skill in the art at the time the invention was made to detect saposin in blood or plasma in the O'Brien method of detecting lysosomal storage disease, since Sano teaches that saposin is found in blood and plasma.

2. Claims 5, 6, 9-12, 19, 20 and 36 are rejected under 35 U.S.C. 103(a) as being unpatentable over O'Brien et al. (1991) Saposin proteins: structure, function, and role in human lysosomal disorders, THE FASEB JOURNAL, vol. 5(3), 301-8, in view of Sano et al., "Sphingolipid hydrolase activator proteins and their precursors", Biochemical and biophysical research communications, 165 (3), pp. 1191-7, 1989, and further in view of Dubensky et al., 6,376,236.

The method of O'Brien in view of Sano discloses the invention as claimed (see above). More specifically, O'Brien discloses the correlation between an accumulation of saposin and Gaucher disease in patients, Gaucher disease being a well known lysosomal disease.

However, neither O'Brien nor Sano specifically teach the step of monitoring the progression of the disease (claim 5), the patient undergoing treatment for the lysosomal storage disorder (claim 6), the patient being an infant (claim 9) or fetus (claim 10), (claim 11), nor the step of determining a treatment program (claims 19 and 20), nor the indication of positive treatment (claims 5, 11, 12 and 36.)

Dubensky discloses a method of treating Gaucher's disease (col. 120, lines 33-59.)

As to claims 5, 6, 11, 12 and 36, it would have been obvious to measure the level of the saposin in a second tissue sample from the patient, the first and second samples being obtained at different times; and comparing the levels in the samples to indicate progression of the disease since Dubensky teaches that some lysosomal storage disorders can be responsive to treatment, thus teaching that after treatment, the disorder can be detected by the disclosed method to determine whether the disorder is responsive to the treatment.

Similarly, as to claim 20, it would have been obvious to determine a treatment program based on the measurement since Dubensky teaches that some lysosomal storage disorders can be responsive to treatment, thus teaching that patients can undergo one of these treatments for lysosomal disorder.

Dubensky further teaches that Gaucher's disease affects infants and fetuses, as well as adults, (col. 120, lines 33-59.) It would have been obvious to one of ordinary skill in the art to use the method taught by O'Brien in view of Sano to detect Gaucher's disease in infants and fetuses, since Dubensky teaches that Gaucher's disease affects infants and fetuses, (claims 9 and 10.) As to claim 19, it would have been obvious to one of ordinary skill in the art to inform the patient or a parent or guardian of an infant of the presence of the lysosomal storage disorder, as would be desirable to allow a patient or parent or guardian to permit treatment of the disease.

3. Claim 16 is rejected under 35 U.S.C. 103(a) as being unpatentable over O'Brien, et al. (1991) Saposin proteins: structure, function, and role in human lysosomal disorders, THE FASEB JOURNAL, vol. 5(3), 301-8, in view of Sano et al., "Sphingolipid hydrolase activator proteins and their precursors", Biochemical and biophysical research communications, 165 (3), pp. 1191-7, 1989, as applied to claims 1 and 15, and further in view of Stastny, J., et al. (1992) Production and Characterization of a Monoclonal Antibody to Human Saposin C, HYBRIDOMA, vol. 11, 351-359.

O'Brien in view of Sano disclose the invention substantially as claimed (see above), except for the antibody being a monoclonal antibody.

Stastny discloses a monoclonal antibody (68-12) that reacts with saposin C. It would have been obvious to use this monoclonal antibody in the method taught by O'Brien in view of Sano in order to detect the level of saposin C because the high specificity of monoclonal antibodies for their corresponding antigen (in this case saposin C) would provide for a more sensitive assay for the detection of saposin C.

Response to Arguments

Applicant still has not overcome the 112 rejections, as further explained above.

Applicant's arguments regarding the 102 and 103 rejections have been considered but are moot in view of the new ground(s) of rejection.

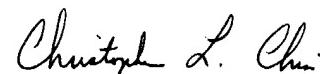
Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Ann Y. Lam whose telephone number is 571-272-0822. The examiner can normally be reached on M-Sat 11-6:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on 571-272-0823. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

A.L.



CHRISTOPHER L. CHIN
PRIMARY EXAMINER
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12/12/04